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US

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Published

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(54) Title: FUNGAL PROMOTERS ACTIVE IN THE PRESENCE OF GLUCOSE

(57) Abstract

(30) Priority data: 932,485

A method is described for the identification and cloning of promoters that express under a defined environmental condition, such as growth in glucose medium. Using this method, five *Trichodermal* promoters capable of the high expression of operably linked coding sequences are identified, one of which is the promoter for *T. reesei tefl*. Also provided are altered *cbh1* promoters, altered so that glucose no longer represses expression from such promoter. The invention further provides vectors and hosts that utilize such promoters, and unique fungal enzyme compositions from such hosts.

CCGCGGACTG CGCATCATGT	1740
TEGTGETCAG TEGGEETGEA	1800
ATGCTCGTCT GGTGGCACTT	1860
GCGCTGGACT CACGCTACGA	1920
GACCCTATGT CCTGACAACG	1980
CGCGTCCACG TACGGAGTTA	2040
GTCTGCGCAG AAGAACGTTG	2100
GGAATTCACC CTGCTTGGCA	2160
AGTGACTTAC CATGAACCCC	2220
TAAGGTGCGG CTTGAACGGA	2280
GCAAGTATCC CACCAACACC	2340
GTCCCCGCGA TCTGAAGTTC	2400
CCAACAACGC AAACACGGGC	2460
GGGAGGCCAA CTCCATCTCC	2520
	2580
GCGATCCCGA TGGCTGCGAC	2640
CTGGCTCAAG CTTTACCCTC	2700
	2760
	2820
	2880
	2940
	3000

FIG.16A

SUBSTITUTE SHEET

AAGCAGCTGA CTGAGATGTT ACAGTACTAC	GCCAACATGC	TGTGGCTGGA	CTCCACCTAC	3060
CCGACAAACG AGACCTCCTC CACACCCGGT	GCCGTGCGCG	GAAGCTGCTC	CACCAGCTCC	3120
GGTGTCCCTG CTCAGGTCGA ATCTCAGTCT	CCCAACGCCA	AGGTCACCTT	CTCCAACATC	3180
AAGTTCGGAC CCATTGGCAG CACCGGCAAC	CCTAGCGGCG	GCAACCCTCC	CGGCGGAAAC	3240
CCGCCTGGCA CCACCACCAC CCGCCGCCCA	GCCACTACCA	CTGGAAGCTC	TCCCGGACCT	3300
ACCCAGTCTC ACTACGGCCA GTGCGGCGGT	ATTGGCTACA	GCGGCCCCAC	GGTCTGCGCC	3360
AGCGGCACAA CTTGCCAGGT CCTGAACCCT	TACTACTCTC	AGTGCCTGTA	AAGCTCCGTG	3420
CGAAAGCCTG ACGCACCGGT AGATTCTTGG	TGAGCCCGTA	TCATGACGGC	GGCGGGAGCT	- 3480
ACATGOCCCC GGGTGATTTA TTTTTTTGT XmaI	ATCTACTTCT	GACCCTTTTC	AAATATACGG	3 540

FIG.16A(Cont.)

```
Title:
                US-10-031-496C-3
RESULT 3
AAQ58015
ID
     AAQ58015 standard; DNA; 1820 BP.
XX
АC
     AAQ58015;
XX
DT
     25-MAR-2003
                  (revised)
DT
     14-SEP-1994
                  (first entry)
XX
     Sequence of plasmid pML017 which carries the shortened form of the
DΕ
     cellobiohydrolase 1 (cbh1) promoter fused to the cbh1 gene.
DE
XX
KW
     Promoter; cellobiohydrolase 1; cbh1; pML017; ss.
XX
OS
     Synthetic.
XX
FH
     Key
                     Location/Qualifiers
FT
     CDS
                     17. .19
FT
                     /*tag= b
FT
                     /label= start codon
FT
                     1773
     misc feature
FT
                     /*tag= a
FΤ
                     /label= KspI-XmaI fragment
FT
                     /note= "contains cbh1 gene"
XX
     WO9404673-A1.
PN
XX
PD
     03-MAR-1994.
XX
ΡF
     19-AUG-1993;
                   93WO-FI000330.
XX
     19-AUG-1992;
PR
                  92US-00932485.
XX
PΑ
     (ALKO-) ALKO OY AB.
XX
PΙ
     Nakari TH, Onnela M,
                           Ilmen MH, Nevalainen KMH, Pentitilae ME;
XX
DR
     WPI; 1994-083192/10.
XX
PT
     Cloning promoters active in a desired environmental condition - used
     partic. for expression of genes in Trichoderma fungal hosts in glucose-
PT
PΤ
     contg. medium.
XX
PS
     Example; Fig 16A; 120pp; English.
XX
CC
     AAQ58015 shows the sequence of the KspI-XmaI fragment that contains the
CC
     chromosomal cbhl gene. pML017 was constructed for the production of CBH1
CC
     on glucose. The plasmid pML016del15(11) was digested with the enzymes
CC
     KspI and XmaI (which is 76 nucleotides downstream from the translation
     stop codon of the cbh1 gene. The vector part contg. the shortened cbh1
CC
CC
     promoter, the cbhl terminator and the pBR322 sequence was ligated to the
CC
     chromosomal cbhl gene isolated as a KspI-XmaI-fragment from the
CC
     chromosomal gene bank of Trichoderma reesei. The sequence of this
CC
     fragment is given in FT. (Updated on 25-MAR-2003 to correct PN field.)
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XX

SQ Sequence 1820 BP; 388 A; 577 C; 478 G; 377 T; 0 U; 0 Other;

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